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	Filing Date		2006-01-18	
	First Named Inventor	BEN-SASSON et al.		
	Art Unit	3762		
	Examiner Name			
	Attorney Docket Number	BEN-SASSON13A		

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1	KELLEHER et al., Characterization of rhodopsin kinase purified from bovine rod outer segments, The Journal of Biological Chemistry, 265(5)2632-2639 (1990)	<input type="checkbox"/>
2	KRUPNICK et al., Arrestin-rhodopsin interaction multi-site binding delineated by peptide inhibition, The Journal of Biological Chemistry, 269(5)3226-3232 (1994)	<input type="checkbox"/>
3	BENOVIC et al., Synthetic peptides of the hamster β 2-adrenoceptor as substrates and inhibitors of the β -adrenoceptor kinase, Er. J. Clin. Pharmac. 30:3S-12S (1990)	<input type="checkbox"/>
4	VARRAULT et al., 5-Hydroxytryptamine1A receptor synthetic peptides mechanisms of adenylyl cyclase inhibition, The Journal of Biological Chemistry, 269(24)16720-16725 (1994)	<input type="checkbox"/>
5	TIMOSSI et al., Structural determinants in the second intracellular loop of the human follicle-stimulating hormone receptor are involved in Gs protein activation, Molecular and Cellular Endocrinology, 189:157-168 (2002)	<input type="checkbox"/>
6	CHUNG et al., NMR structure of the second intracellular loop of the α 2A adrenergic receptor: Evidence for a novel cytoplasmic helix, Biochemistry, 41:3596-3604 (2002)	<input type="checkbox"/>
7	JIMENEZ-CERVANTES et al., The Pro162 variant is a loss-of-function mutation of the human melanocortin 1 receptor gene, The Journal of Investigative Dermatology, 117(1)156-158 (2001)	<input type="checkbox"/>
8	LEE et al., Vascular endothelial cell adherens junction assembly and morphogenesis induced by sphingosine-1-phosphate, Cell, 99:301-312 (1999)	<input type="checkbox"/>
9	OKADA et al., Activation of rhodopsin: new insights from structural and biochemical studies, TRENDS in Biochemical Sciences, 26(5)318-324 (2001)	<input type="checkbox"/>
10	BALLESTEROS et al., G protein-coupled receptor drug discovery: Implications from the crystal structure of rhodopsin, Current Opinion in Drug Discovery and Development, Current Drugs, 4(5)561-574 (2001)	<input type="checkbox"/>

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